WASTER: Practical *de novo* phylogenomics from low-coverage short reads

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Abstract

The advent of affordable whole-genome sequencing has spurred numerous large-scale projects aimed at inferring the tree of life, yet achieving a complete species-level phylogeny remains a distant goal due to significant costs and computational demands. Traditional species tree inference methods, though effective, are hampered by the need for high-coverage sequencing, high-quality genomic alignments, and extensive computational resources. To address these challenges, this study introduces WASTER, a novel *de novo* tool for inferring species trees directly from short-read sequences. WASTER employs a k-mer based approach for identifying variable sites, circumventing the need for genome assembly and alignment. Using simulations, we demonstrate that WASTER achieves accuracy comparable to that of traditional alignment-based methods, even for low sequencing depth, and has substantially higher accuracy than other alignment-free methods. We validate WASTER's efficacy on real data, where it accurately reconstructs phylogenies of eukaryotic species with as low depth as 1.5X. WASTER provides a fast and efficient solution for phylogeny estimation in cases where genome assembly and/or alignment may bias analyses or is challenging, for example due to low sequencing depth. It also provides a method for generating guide trees for tree-based alignment algorithms. WASTER's ability to accurately estimate trees from low-coverage sequencing data without relying on assembly and alignment will lead to substantially reduced sequencing and computational costs in phylogenomic projects.

Keywords: Alignment free, Assembly free, Low coverage reads, Phylogenomics, WASTER

1 Introduction

Whole-genome sequencing has recently replaced single markers sequencing as the dominant method for estimating phylogenies, and has been the focus of many recent large-scale efforts for a broad set of species [1–10].Nevertheless, these efforts are still short of realizing complete species-level phylogenies for most clades. Major obstacles include the costs of high-coverage sequencing, extensive human effort in running species tree inference pipelines, and substantial computational resource demanded [11]. In addition, alignment challenges may possibly affect the results in downstream phylogenetic analyses [12].

Traditional species tree inference pipelines typically involve several key steps: assembling reads into genomes, aligning orthologous genomic sequences, optionally reconstructing gene trees from aligned genomes, and inferring the species tree from gene trees or directly from the aligned genomes (Fig. 1a). The recent development of CASTER significantly speeds up accurate species tree inference in the presence of incomplete lineage sorting (ILS), directly from aligned genomes [13]. CASTER mitigates the computational bottleneck from aligned genomes to species tree. However, the necessity for high coverage sequencing persists, with traditional pipelines usually requiring coverage above 30X for reliable genome assemblies, thus imposing financial and computational challenges [14, 15]. Additionally, Progressive Cactus, the leading tool for aligning a large number of genomes, relies on a user-specified guide-tree [16]. The use of a guide-tree for alignment may potentially lead to circular inferences when subsequently estimating the tree using the very same alignment. These factors underscore the need for accurate alignment-free and assembly-free species tree inference methods.

Several assembly-free methods have been proposed [17–21], but few are suitable for inferring trees from reads covering entire eukaryotic nuclear genomes. Some methods rely on mapping reads to reference sequences, thus requiring alignments of annotated orthologous sequences as input [21]. This limits their applicability, especially in the absence of closely related reference genomes. Others compute pairwise evolutionary distances using k-mers, invoking distance-based methods to infer species trees [17–20]. However, these methods struggle to infer accurate phylogenies for challenging trees with short internal branches.

In this article, we introduce WASTER (Without Alignment/Assembly Species Tree EstimatoR), a novel *de novo* tool that infers species trees directly from short reads. WASTER operates by initially calling variable sites using k-mers from raw sequencing reads and then employs CASTER-site to infer the species tree from the matrix of variable sites (Fig. 1a). We evaluate the method extensively using simulations and applications to real data and show that it substantially outperforms other alignment-free methods and is as accurate as state-of-the-art alignment-based methods, even when assuming that the genome assembly and alignment are error free.

2 Results

2.1 WASTER: k-mer-based variable site identification

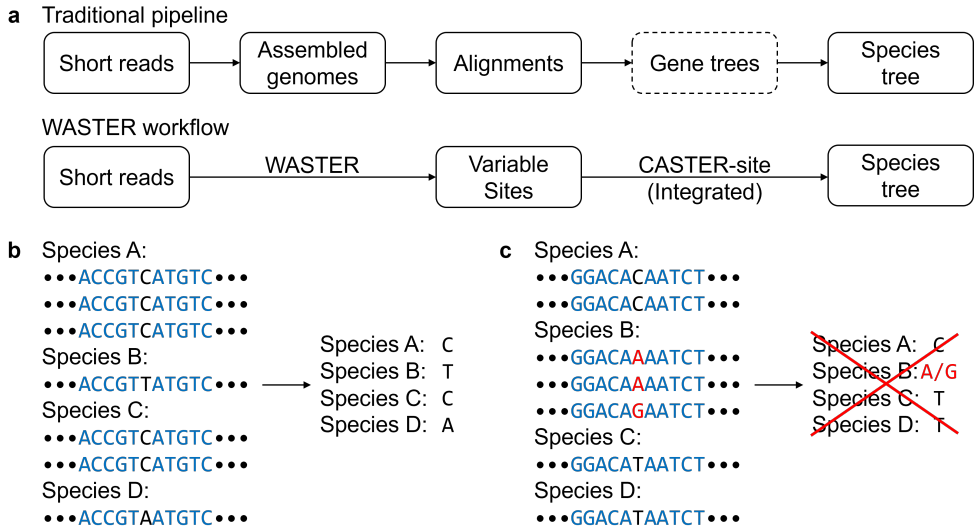

Fig. 1 WASTER workflow. a, Overview of the WASTER workflow compared to the traditional species tree inference pipeline. **b**, WASTER indentifies variable sites by detecting k-mers that differ only at the central base. **c**, If two variants of a k-mer are found within the same species, WASTER removes the entire aligned site.

The variable site identification mechanism within WASTER involves a k-mer based approach. Initially, WASTER decomposes input reads or genomes into k-mers of a specified length k, where k is an odd number (19 by default). The process then groups these k-mers by matching all positions except the central one. For a given group of k-mers, if the k-mers from each species are identical, including the central position, then the central position of this k-mer group is designated as one variable site (Fig. 1b). In contrast, if any two k-mers from the same species differ, the entire group is disregarded, reducing the possibility of paralogy, false homology, or sequencing errors. This approach, however, results in the exclusion of heterozygous sites (Fig. 1c). Following the variable site identification, the concatenated variable sites are utilized as input for CASTER-site, which undertakes the inference of the species tree. Due to WASTER's inability to distinguish between the forward and reverse strands, it employs the T92 substitution model [22], a simplified model that does not account for strand bias.

2.2 Evaluation in simulations

In the main text, we demonstrate the performance of WASTER on one simulated dataset. Additional benchmarking results on other simulated datasets are provided in the supplement (see Figs. A1–A3).

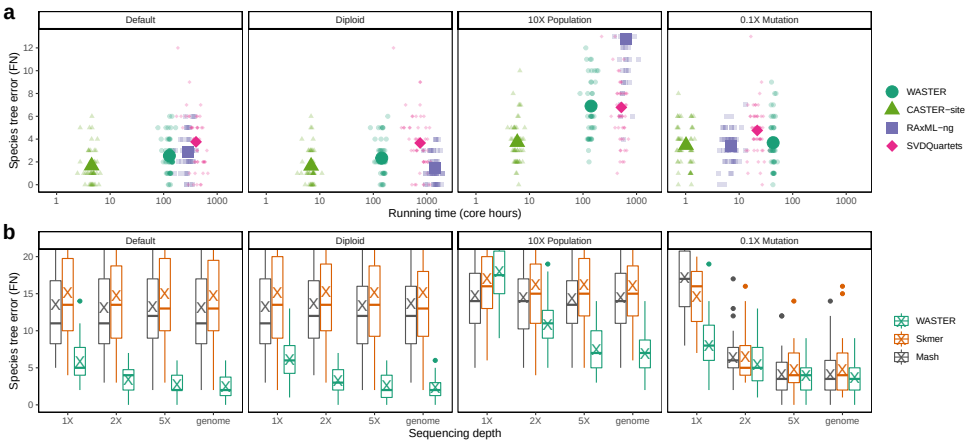

Fig. 2 Benchmarking WASTER via simulations. All simulations are performed on the 201-species phylogeny described in [13]. **a**, The number of branches differing between the inferred and true species trees (bipartition false negatives) versus running time for WASTER and three alignment-based species tree inference methods using aligned genomes under various settings. Small symbols represent individual replicates, while large, opaque symbols denote mean values across 50 replicates. **b**, The number of branches differing between inferred and true species trees (FN) versus sequencing depth for WASTER, Mash, and Skmer on simulated short reads under different settings. Inferred species trees using true genomes are included as control groups. Under 0.1X mutation rate setting, Skmer fails to output phylogenies on 36/24/25/21 replicates for 1X/2X/5X/genome cases, respectively.

2.2.1 Simulated alignments

We first evaluated the performance of WASTER versus alignment-based methods using the SR201 dataset which comprised simulated alignments from 201 species [13]. This dataset incorporates many of the intricacies of molecular evolution. To avoid unrealistic strict molecular clocks, this dataset allows variation in generation time and population size across different branches of the species tree. Simulations were conducted under the Hudson model [23] for recombination and the GTR substitution model [24], with parameters adjusted for varying mutation rates across the genome. Notably, the equilibrium nucleotide frequencies differ from the no strand-bias model assumed by WASTER. We juxtaposed WASTER with several leading alignment-based methods:

1. RAxML-ng [25], a maximum-likelihood approach, widely used for reconstructing concatenated sequence tree yet statistically inconsistent,
2. SVDQuartets [26], an invariant-based method maintaining statistical consistency under ILS,
3. CASTER-site [13].

Although providing true simulated alignments as inputs, rather than assembled genomes from simulated reads, disproportionately favored alignment-based methods, WASTER still demonstrated accuracy on par with them. Across all experiments, WASTER's species tree branch estimation error averaged at 3.9 branches, significantly lower ($p < 10^{-5}$) than RAxML-ng (5.1 branches) and SVDQuartets (4.7 branches),

but higher ($p < 10^{-15}$) than CASTER-site (2.6 branches). Moreover, WASTER's total runtime was comparable to the time required for species tree estimation from alignments alone in traditional workflows, being 5.1 times faster than RAxML-ng and 3.7 times faster than SVDQuartets (Fig. 2a).

We also explored WASTER's performance under various settings, contrasting it with alignment-based methods under different ploidies, ILS levels, and mutation rates. Replacing haploid genomes with diploid genomes did not significantly impact ($p \approx 0.11$) WASTER's accuracy, mirroring results seen with CASTER-site and SVDQuartets. RAxML-ng, employing a diploid model, improved accuracy at the cost of increased runtime. An increase in effective population size by 10X to model elevated ILS levels notably diminished the accuracy of all methods, including WASTER ($p < 10^{-15}$). Here, RAxML-ng's performance was notably poorer than WASTER and the other methods. To simulate less genetic divergence among species, we reduced the mutation rate to 10%, resulting in a statistically significant decrease in accuracy for all methods ($p < 0.05$), attributed to a reduction in informative sites. Under this setting, the runtime of alignment-based methods improved substantially compared to the default setting. However, due to bottlenecks in the variable site identification step, WASTER's runtime improvement was less pronounced (Fig. 2a).

2.2.2 Simulated short reads

We extended our analysis to assess WASTER's performance utilizing Illumina-like short reads, generated from the SR201 dataset alignments using the ART simulator [27]. This evaluation focused on the impact of varying coverage levels on WASTER's accuracy. Our findings indicated a direct correlation between increased coverage and enhanced accuracy of WASTER across all settings. A notable observation was that at 5X coverage, the species tree error for WASTER, based on short reads, averaged 4.2 branches, deteriorating by less than 10% compared to the error obtained using true simulated genomes.

Furthermore, we contrasted WASTER's accuracy with that of Mash [17] and Skmer [19], two frequently used k-mer-based assembly-free and reference-free pairwise evolutionary distance estimators. The k-mer-based distance methods, commonly employed for phylogenetic reconstruction by providing distance matrices to FastME [28], demonstrated consistent species tree errors of approximately 15 branches across all but 0.1 mutation rate settings, irrespective of coverage level. This is likely attributable to their lack of robustness to high evolutionary distances, where fewer k-mers are shared across species, underscoring the superiority of WASTER.

2.3 Applications to raw reads

We also compared phylogenies estimated using WASTER, Mash, and Skmer from low-coverage short reads, to phylogenies constructed using traditional methods employing high-coverage reads. Our analyses span multiple taxonomic groups:

1. **Rats:** We evaluated 18 rat species, particularly focusing on the *Niviventer* genus [29]. The established phylogeny, derived from 14X coverage reads (approximately 39 Gbps), was obtained using MrBayes [30] (concatenation) and ASTRAL [31], both

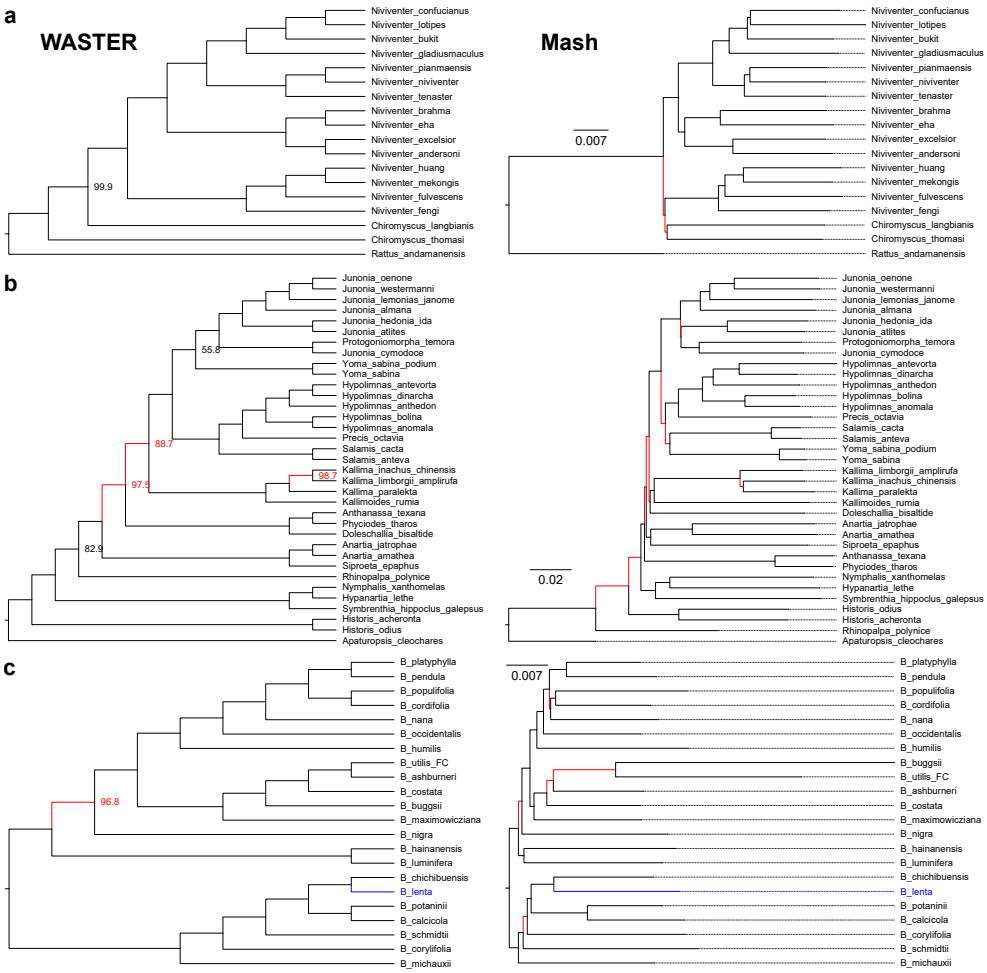

Fig. 3 WASTER and Mash on low-coverage short reads. Phylogenies reconstructed using (**a**) 1.5X WGS reads of rat species, (**b**) 2 Gbp butterfly WGS reads, and (**c**) 0.3–3 Gbp birch RAD-seq reads. WASTER branch supports (%) are shown next to the branches, and 100% supports are omitted. Branches that are incongruent with published phylogenies based on high-coverage reads using traditional pipelines are colored red. B. lenta is placed differently from the published phylogeny due to introgression. See also Fig. A4.

yielding identical results. We downsampled the reads to 1.5X coverage (3.75 Gbps) for reconstructing phylogenies with WASTER, Mash, and Skmer. The phylogeny reconstructed by WASTER matched the established one, while the phylogenies by Mash and Skmer differed from the established phylogeny by two and five branches, respectively (Fig. 3a and A4a).

2. **Butterflies:** This study involved 35 butterfly species from the Nymphalidae family [32], with a published phylogeny derived from RAxML on genome-wide variable sites. We standardized the downsampled reads to 2 Gbps across species. WASTER's reconstruction deviated from the published phylogeny by three branches. Notably,

the phylogeny observed within the *Kallima* genus aligns with the phylogeny inferred by CASTER-site, suggesting potential introgression [13]. The discrepancies in the deeper splits could potentially be attributed to rapid radiation causing short internal branch lengths associated with high statistical uncertainty [32]. Mash and Skmer, however, differed by 8 and 9 branches, respectively (Fig. 3b and A4b).

3. **Birches:** We analyzed high-coverage RAD sequences (3 Gbps on average) for *Betula utilis* FC and *Betula buggsii* [33] and low-coverage RAD sequences (0.3 Gbps on average) for 20 other birch species [34]. The established phylogeny, inferred using RAxML on aligned orthologous internal transcribed spacer (ITS) sequences, was compared with our reconstructions. WASTER, Mash, and Skmer all placed *B. lenta* as the sister of *B. chichibuensis*, rather than the sister of *B. michauxii*, likely due to introgression (Fig. A5ab), which is confirmed using Patterson's D-statistics [35] (Fig. A5c). Beside *B. lenta*, the WASTER inferred phylogeny differed from the published phylogeny by one branch, while Mash and Skmer differed by five and seven branches, respectively (Fig. 3c and A4c). Noticeably, Mash misgrouped the two species with high coverage as sister taxa, suggesting a potential bias in the method under conditions of non-uniform coverage.
4. **Lizards:** We conducted a similar analysis on 11 Anolis lizard species [36]. The original phylogeny was based on 9X coverage reads. ExaML [37] (concatenation) and ASTRAL generated consistent phylogenies. Upon downsampling to 1.5X coverage (2.25 Gbps), the reconstructed phylogenies by WASTER and Mash matched the published one, while Skmer's reconstruction deviated by one branch (Fig. A4d).
5. **Mandarin fishes:** Our analysis included three mandarin fish species and one outgroup [38], with *Siniperca chuatsi* sequenced using high-coverage long reads and the others with approximately 40X coverage short reads. The published phylogeny was deduced using RAxML [39]. Using downsampled short reads with 2X coverage (1.5 Gbps), as well as the assembled *Siniperca chuatsi* genome, all three methods successfully reconstructed the published phylogeny (Fig. A4e).
6. **Mushrooms:** We examined short reads from four species in the *Tricholomopsis* genus and one outgroup [40]. The established phylogeny, inferred using RAxML on 450 single-copy orthologous genes, was compared with our phylogenies using downsampled reads (1 Gbps). WASTER and Mash successfully reconstructed the published phylogeny, but Skmer did not (Fig. A4f).

These results demonstrate that phylogenies reconstructed using WASTER on low-coverage reads closely align with those derived from traditional pipelines from high-coverage reads, significantly outperforming Mash and, even more so, Skmer in accuracy.

2.4 Applications to assembled genomes

In addition to inferring phylogenies of closely related species, WASTER can also generate guide trees for more distantly related species with sufficient accuracy, particularly beneficial when using Progressive Cactus. We demonstrate this with guide trees created by WASTER, Mash, and Skmer for assembled genomes of species in the Primates and Passeriformes orders:

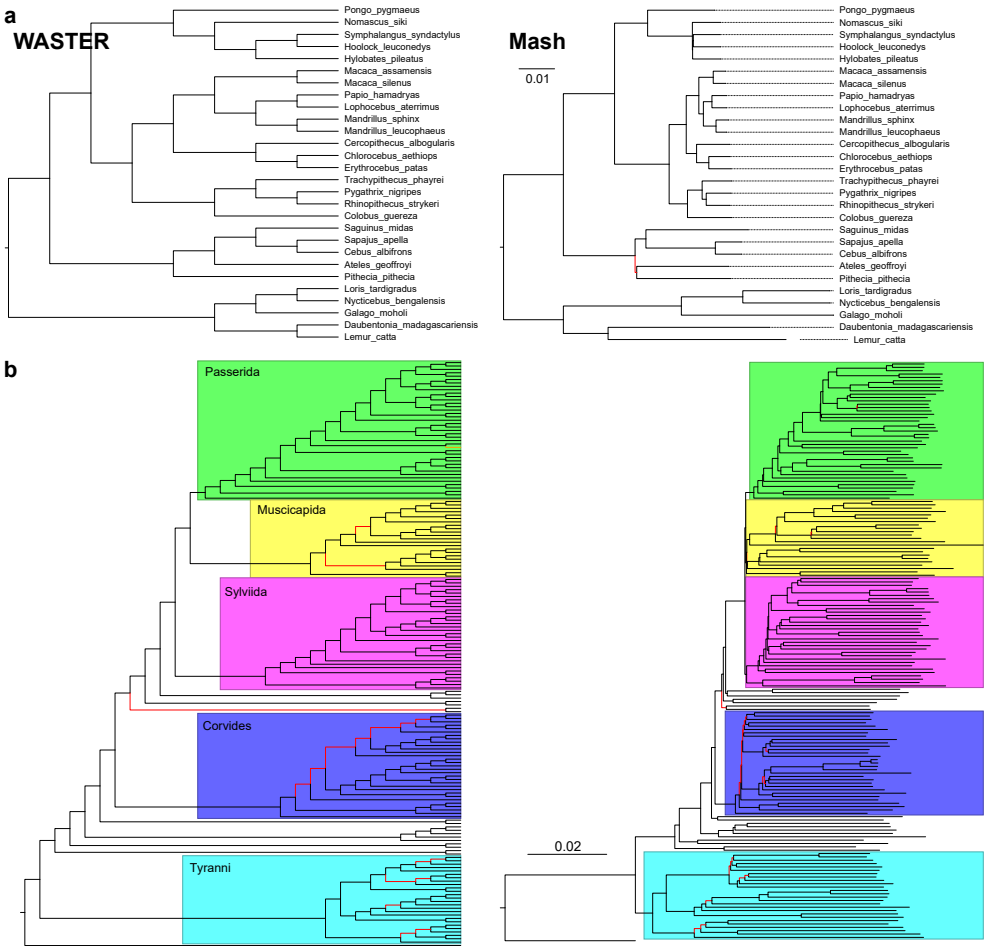

Fig. 4 WASTER and Mash on assembled genomes. a, Phylogenies reconstructed using 28 assembled primate genomes. WASTER provides 100% supports for all branches. **b**, Phylogenies reconstructed from assembled genomes of 173 perching birds, with species names and support values omitted. Branches incongruent with the published phylogenies are highlighted red. See also Fig. A6.

1. **Primates:** Using a dataset of 28 assembled primate genomes [41], WASTER accurately recovered the published phylogeny reconstructed by ExaML on aligned genomes. However, the reconstructions by Mash and Skmer deviated by one and two branches, respectively (Fig. 4a and A6a).
2. **Perching birds:** The dataset of 173 perching bird genomes [10], well-known for their challenging phylogeny, was analyzed. The published phylogeny, reconstructed by ASTRAL+RAxML on aligned genomes, was compared with our reconstructions. WASTER and Mash successfully recovered major clades of Passeriformes (Passerida, Muscicapida, Sylviida, Corvides, and Tyranni), differing from the published phylogeny by 17 and 24 branches, respectively (Fig. 4b). Skmer, however,

failed to recover 64 out of 170 branches in the published phylogeny, including misplacing the cinnamon ibon (Hypocryptadius cinnamomeus) among Sylviida, instead of Passerida (Fig. A6b).

3 Discussion

The increasing availability of whole-genome sequencing data has not resulted in a corresponding increase in phylogenetic inference using aligned whole genomes. A major impediment has been the extensive computational resources and significant human effort required, rendering such approaches impractical for many researchers. WASTER addresses this challenge by offering a computationally efficient and automated solution, facilitating broader adoption of whole-genome phylogenetic analyses. Additionally, WASTER's output can reduce the cost of many projects by providing high-quality phylogenies using assembly-free low-coverage sequencing, thereby greatly reducing the cost of phylogenomics. This type of application will enable whole genome approaches to be adopted by a much larger share of the taxonomic community that does not have access to the type of resources necessary to produce high-quality assemblies for all species of interest. Even when assembly is possible, WASTER can play a crucial role in verifying the accuracy of the phylogenetic inferences that rely on many steps of assembly and alignment subject to potential human errors. Furthermore, WASTER can be used to produce guide trees for progressive alignment.

WASTER also presents several avenues for future development and refinement. For example, there exists a promising potential for adapting WASTER to identify possible gene-flow events, including ancient ones, using phylogenetic asymmetries and for inferring phylogenetic networks. Another key area for advancement is to extend WASTER to estimate branch lengths in presence of ILS and, potentially, gene-flow. Finally, integrating WASTER with other variable-site-based tools would create a more comprehensive and user-friendly platform, broadening the scope of assembly-free phylogenomic and evolutionary studies.

4 Methods

4.1 WASTER

WASTER initially identifies aligned variable sites using Algorithm 1, described below, before leveraging CASTER-site for species tree inference. Acknowledging its limitation in distinguishing between forward and reverse DNA strands, WASTER operates under the assumption of no strand-bias, equating the equilibrium frequencies of A to T and C to G. Consequently, differently from the conventional CASTER-site that assumes the F84 model, WASTER operates under the T92 model. This model can be easily reduced from F84 model by assuming equal equilibrium frequencies of complementary nucleotides (Fig. 5).

Algorithm 1 WASTER's algorithm to indentify a set of aligned variable sites S using sets of k-mers $K_1, \ldots, K_n$, where each set represents all k-mers in the input reads/genomes of a species.

```
procedure AddKmer(M, i, s)                    ▷ M will be modified in this procedure
    L, c, R ← the first (k−1)/2-mer, the central 1-mer, and the last (k−1)/2-mer of s
    j ← the (k − 1)-mer combining L and R
    if M[i, j] ≠ "-" and M[i, j] ≠ c then
        M[i, j] ← "N"                                          ▷ Marked as inconsistent
    else
        M[i, j] ← c
    end if
end procedure
procedure WASTER
    S ← empty alignment, M ← empty look-up table              ▷ a map or a dictionary
    for i in {1, ..., n} do
        for j in all (k − 1)-mers do
            M[i, j] ← "-"                                      ▷ M is initialized with all gaps
        end for
        for s in K_i do
            AddKmer(M, i, s)
            AddKmer(M, i, reverse complement of s)
        end for
    end for
    for j in all (k − 1)-mers do
        if M[·, j] doesn't contain "N" and M[·, j] is not invariant then
            Add M[·, j] as one column of S
        end if
    end for
end procedure
```

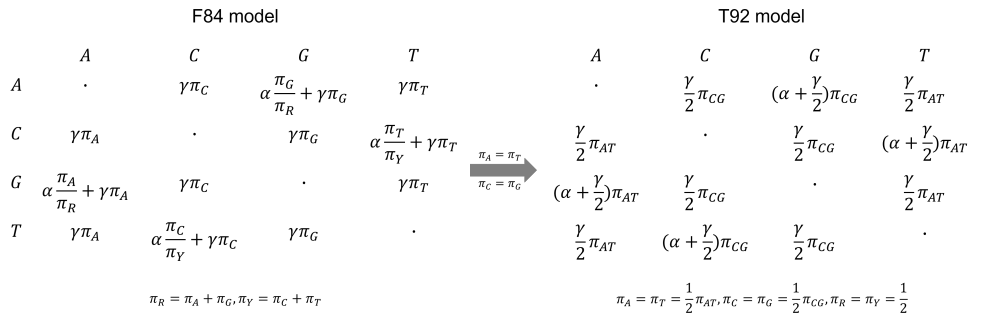

Fig. 5 No-strand bias model. By assuming equal equilibrium frequencies of complementary nucleotides, the F84 model reduces to the T92 model.

4.2 Simulating low-coverage reads

Reads with 1X, 2X, and 5X coverage in the simulation studies were simulated using ART Illumina v2.5.8 with the following command:

```
for i in 1 2 5; do
    art_illumina -ss HS25 -sam -i $GENOME -p -l 150 -f $i \
        -m 800 -s 20 -o ${i}X_READS
    cat ${i}X_READS{1,2}.fq > ${i}X_READS_COMBINED.fq
done
```

Each input file contains one FASTA sequence representing a haploid genome or two FASTA sequences representing a diploid genome. Each output file contains reads from both ends.

4.3 Species tree inference

We inferred species tree from simulated and real data using WASTER, Mash, and Skmer. Aside from those alignment free methods, we also reuse inferred species trees by CASTER-site, RAxML-ng, and SVDQuartets on simulated true alignments from a previous study [13] for comparison (Fig. 2a).

4.3.1 WASTER

The following command was used to infer species trees:

```
waster-site -C -t 16 -i $INPUT -o $OUTPUT
```

The input file here is a list of species names followed by the corresponding input files:

```
SpeciesName1    Species1.fq
SpeciesName2    Species2.fq
...
```

4.3.2 Mash

The following script was used to infer species trees:

```
mash triangle -s 1000000 $INPUT > mash.tsv
python3 mashFormatFastme.py mash.tsv mash.phy
fastme -i mash.phy -o $OUTPUT
```

The python script "mashFormatFastme.py" was used to format the output distance matrix by Mash as the input format of FastME:

```
import sys
with open(sys.argv[2], "w") as f:
    lines = []
    for line in open(sys.argv[1]):
        lines.append(line.split())
    n = int(lines[0][0])
    f.write(lines[0][0] + "\n")
    for i in range(1, n+1):
        f.write("\t".join([lines[i][0].split(".")[0]]
        + [lines[i][j] for j in range(1, i)] + ["0"]
        + [lines[j][i] for j in range(i+1, n+1)]) + "\n")
```

4.3.3 Skmer

The following script was used to infer species trees:

```
skmer reference $INPUT -p 16 -t #generating ref-dist-mat.txt
bash skmerFormatFastme.bash ref-dist-mat.txt skmer.phy
fastme -i skmer.phy -o $OUTPUT
```

the python script "skmerFormatFastme.bash" was used to format the output distance matrix by Skmer as the input format of FastME:

```
tail -n +2 $1 | wc -l > $2
tail -n +2 $1 >> $2
sed -i "s/\t/--/g" $2
```

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Appendix A Supplementary figures

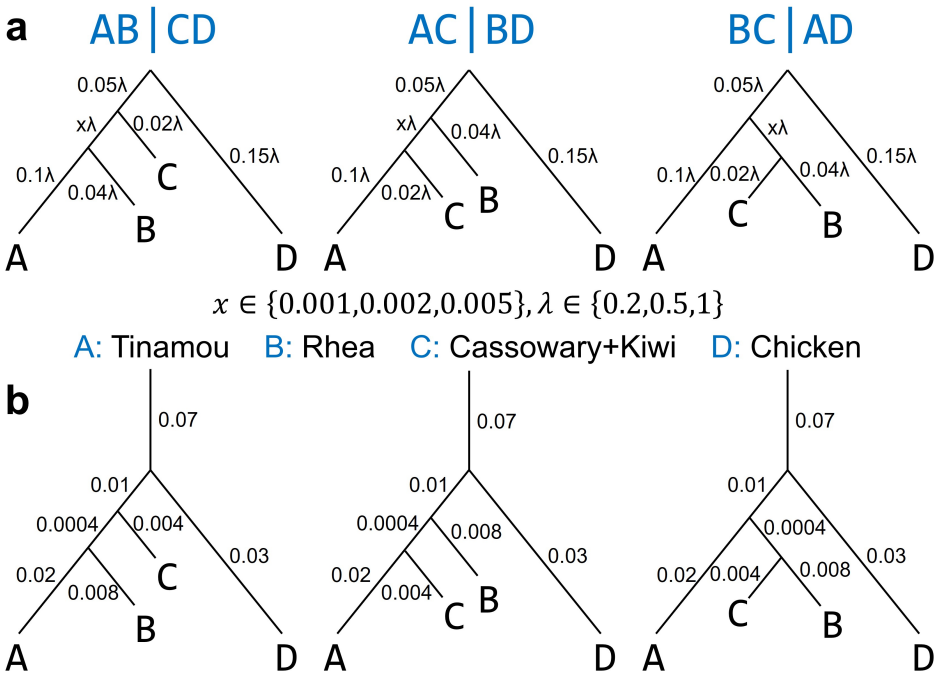

Fig. A1 Experimental setup of additional simulation studies. (**a**) Quartet dataset. We vary tree depth (λ) and internal branch length (x) to create different model conditions. For each condition, we simulate aligned genomes from three topologies (100 replicates per topology), corresponding to Felsenstein's ($AB|CD$ and $AC|BD$) and Farris's zone ($BC|AD$). The branch lengths, labelled in substitution units, are modeled after ratite phylogenies. Aligned genomes are simulated with recombination for various lengths (1 – 20 Mbps) under Hudson model using MSPrime. In all conditions, we maintain a consistent level of ILS by fixing the discordance between the species tree and the local (gene) trees at 60%. Short reads of various coverage levels (1 – 5X) are also simulated using ART. (**b**) GDL dataset. In this dataset, we simulate gene families under various gene duplication and loss (duploss) rate, fixing $x = 0.002$ and $\lambda = 0.2$. Gene duplications and losses are simulated beginning 0.07 substitution unit above the root to allow the emergence of superfamilies. We vary gene duplication rate ($\{0,1,2,4,8\}$X) while maintaining a 1:1 ratio between gene duplication and gene loss rates. For each replicate, we simulate 10,000 gene family trees using Simphy, followed by recombination-free alignments of lengths between 100 and 10,000 bps generated with INDELible.

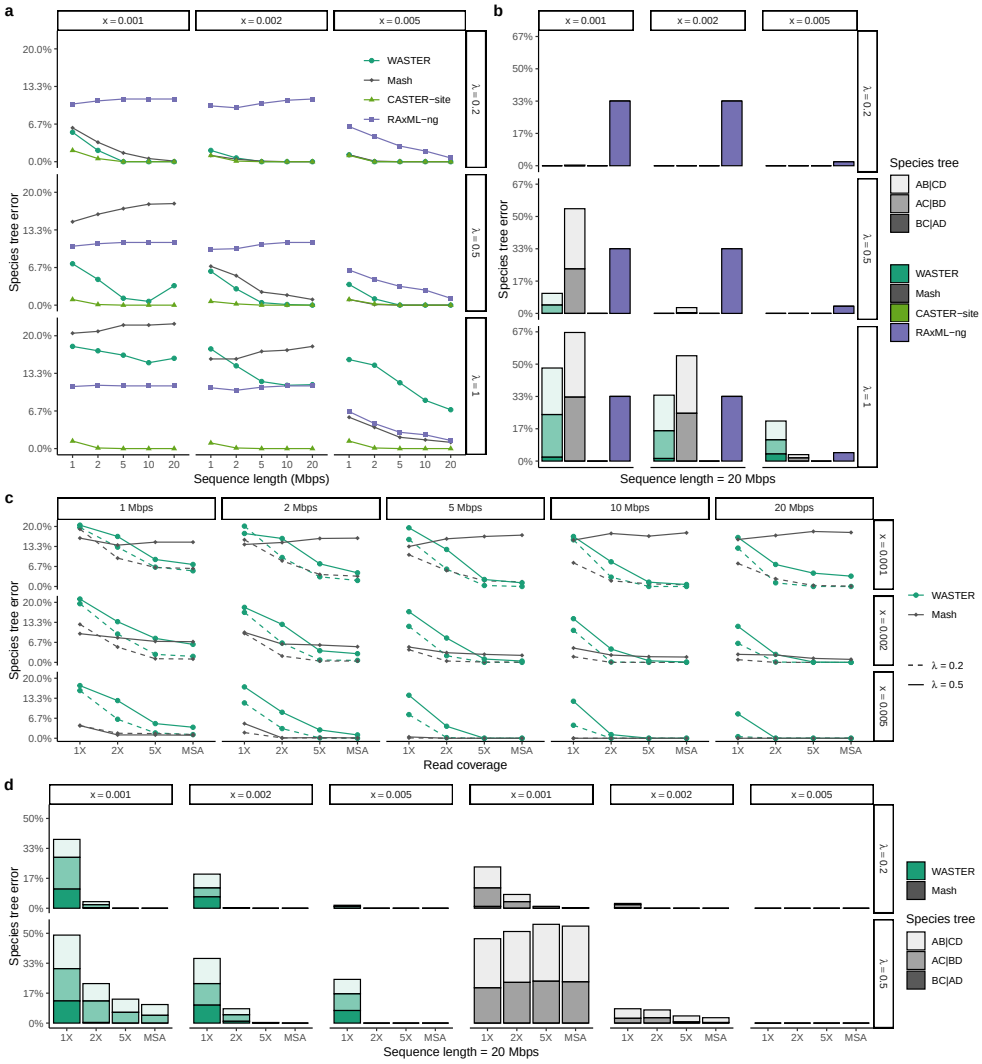

Fig. A2 Benchmarking via Quartet dataset. Species tree estimation error of alignment-free and alignment-based methods on true MSAs by (**a**) sequence length and (**b**) true species tree topology. CASTER-site outperforms all other methods across all conditions. RAxML-ng exhibits long-branch repulsion in the Farris zone tree ($BC|AD$), particularly when the internal branch length is short ($x \leq 0.002$). Both alignment-free methods experience a significant increase in species tree estimation error under deeper phylogenies ($\lambda = 1$). Under shallow phylogenies ($\lambda \leq 0.5$), WASTER outperforms Mash when internal branch length is short ($x \leq 0.002$), although it is also more sensitive to shorter sequences (≤ 2 Mbps). Species tree estimation error of assembly-free methods on simulated short reads and true MSAs by (**c**) sequence length and (**d**) true species tree topology under shallow phylogenies ($\lambda \leq 0.5$). Generally, WASTER performs better at higher coverage (5X), while Mash outperforms WASTER at very low coverage (1X).

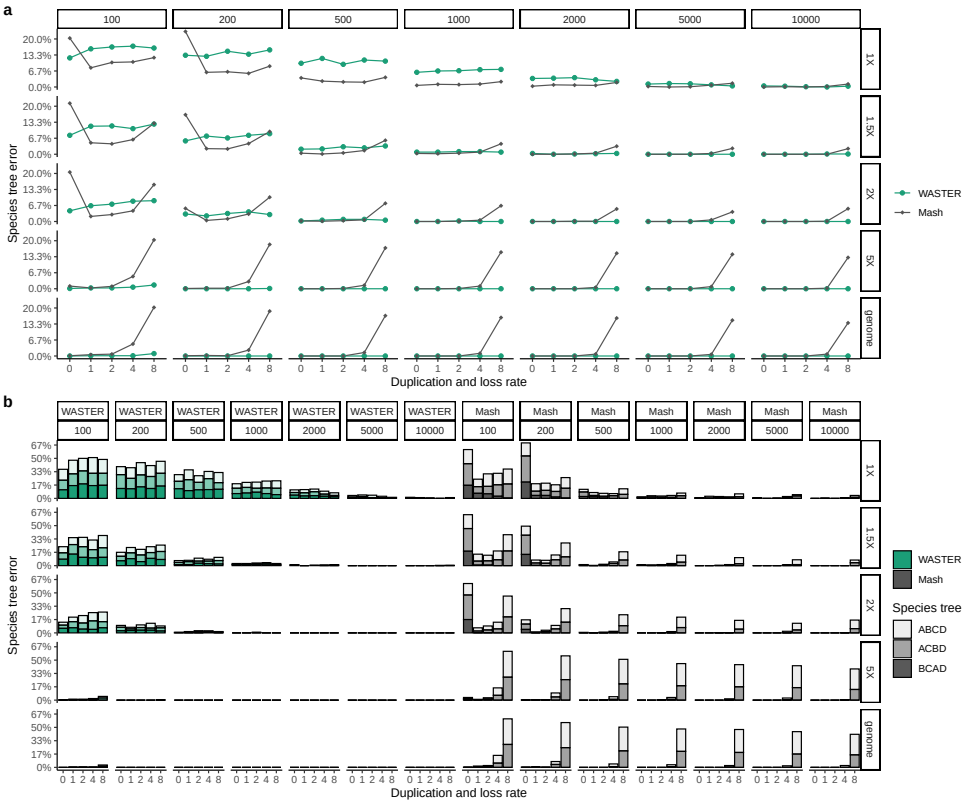

Fig. A3 Benchmarking via GDL dataset. Species tree estimation error of WASTER and Mash on simulated short reads and true MSAs across (**a**) gene duplication and loss (duploss) rates and (**b**) duploss rates and true species tree topology under various sequencing depths (rows) and gene lengths (columns). As genome length increases, WASTER's species tree inference error rate approaches 0%, regardless of duploss rates and sequence depths. In contrast, Mash's error rate rises significantly with increasing duploss rates. Furthermore, at high duploss rates (8X), Mash's error rate also increases with higher sequencing depths.

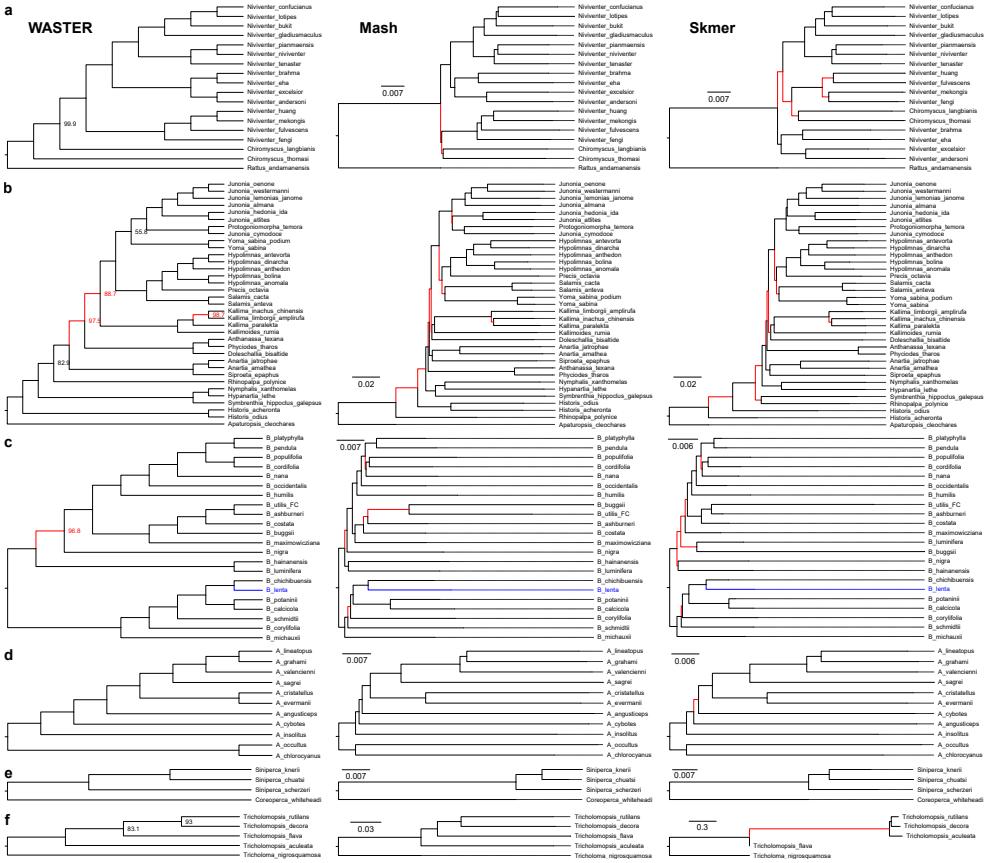

Fig. A4 WASTER, Mash, and Skmer on low-coverage short reads. Phylogenies reconstructed using (**a**) 1.5X WGS reads of rat species, (**b**) 2Gbp butterfly WGS reads, (**c**) 0.3-3Gbp birch RAD reads, (**d**) 1.5X lizard WGS reads, and (**e**) 1.5Gbp mandarin fish WGS reads, and (**f**) 1Gbp mushroom WGS reads. WASTER branch supports (%) are shown next to the branches, and 100% supports are omitted. Branches that are incongruent with published phylogenies based on high-coverage reads using traditional pipelines are colored red. B. lenta is placed differently from the published phylogeny due to introgression.

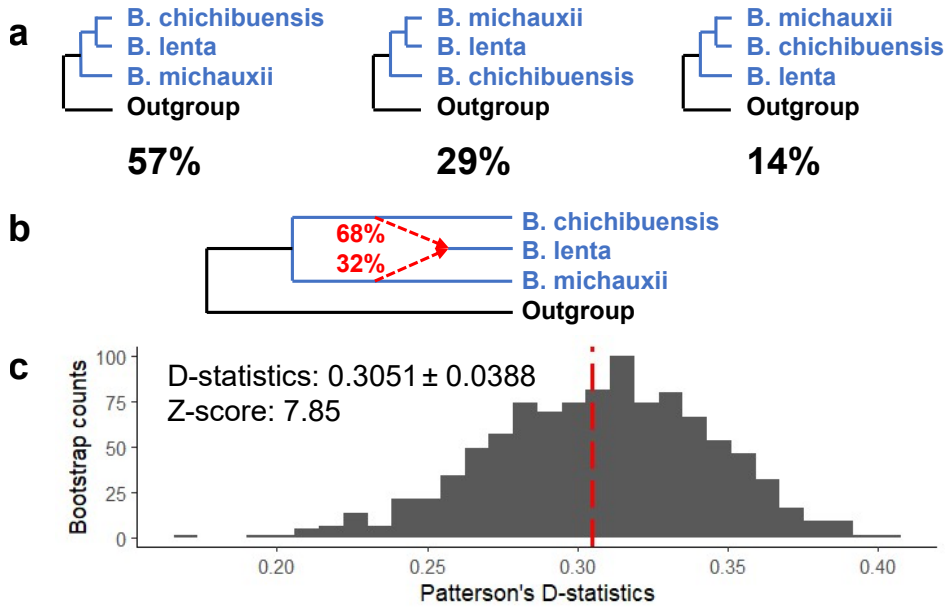

Fig. A5 Non-ILS-like signal around B. lenta. a, Normalized WASTER scores for all quartet topologies with species B. chichibuensis, B. lenta, B. michauxii, and B. maximowicziana (outgroup). Notably, in the absence of introgression, the second and third topologies would be expected to have similar WASTER scores. **b**, One plausible explanation for the observed disparity in WASTER scores is hybrid speciation. In this scenario, 68% of B. lenta genome is derived from B. chichibuensis and 32% originates from B. michauxii. **c**, A histogram of Patterson's D-statistics on 1,000 block bootstrap replicates confirms a non-ILS-like signal with 100% bootstrap support. The D-statistics is based on 4,666 alignments of RAD sequences and the species tree topology (((B. chichibuensis, B. lenta), B. michauxii), Outgroup). The dashed line represents the D-statistics of 0.3051 without bootstrapping. The standard deviation based on 1,000 bootstraps is 0.0388, yielding a Z-value of 7.85.

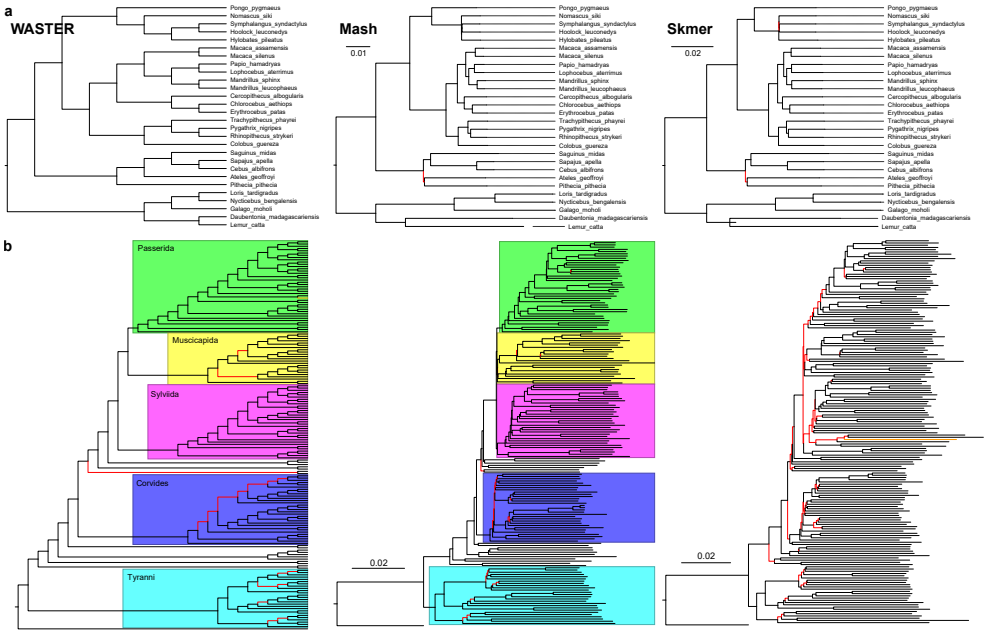

Fig. A6 WASTER, Mash, and Skmer on assembled genomes. a, Phylogenies reconstructed using 28 assembled primate genomes. **b**, Phylogenies reconstructed using assembled genomes of 173 perching birds, with species names and support values omitted. The terminal branch corresponding to cinnamon ibon (Hypocryptadius cinnamomeus) is colored orange. Branches incongruent with the published phylogenies are highlighted red.

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